IN THE CLAIMS:

- 1. (Amended) A method of determining the *initial dose* of a *vitamin D* compound[,] for the treatment of secondary hyperparathyroidism and renal osteodystrophy without increasing the incidence of hypercalcemia comprising:
 - a. measuring a patient baseline PTH (bPTH) value,
 - b. determining [the] a final dose of the vitamin D compound, where the final dose is that dose associated with a first stable clinically significant reduction in patient intact parathyroid hormone (PTH) for the vitamin D compound,
 - applying the baseline PTH value and final dose to regression analysis, and
 - d. calculating the *initial dose* of the *vitamin D compound* from the regression analysis of step c.
- 2. (Original) The method of claim 1 wherein the [linear model] <u>regression</u> <u>analysis</u> is a zero intercept linear model.
- 3. (Original) The method of claim 1 wherein the vitamin D compound is a vitamin D_2 compound.
- 4. (Original) The method of claim 3 wherein the vitamin D_2 compound is paricalcitol.
- 5. (AMENDED) The method of claim 4 wherein the initial dose is <u>patient</u> <u>baseline PTH/80 (bPTH/80)</u>.
- 6. (Amended) [The] A method of treating secondary hyperparathyroidism and renal dystrophy using a vitamin D compound without increasing the incidence of hypercalcemia [claim 1 further] comprising
 - a) measuring a patient baseline PTH value;
- b) determining a final dose of the vitamin D compound associated with a first stable clinically significant reduction in patient PTH for the vitamin D compound;
 - c) applying the baseline PTH and final dose to regression analysis;
- d) calculating the initial dose of the vitamin D compound from the regression analysis of step c; and
- e) [administration of] <u>administering</u> the initial dose determined <u>in step d</u> to the patient.
- 7. (Amended) A method of treating elevated <u>intact parathyroid hormone</u> (PTH) in a patient commencing treatment for [ESRD] <u>end stage renal disease</u>, the method comprising:

- a. determining the initial dose of a vitamin D compound from a regression analysis based on a patient baseline PTH (bPTH) and a final dose of the vitamin D compound associated with a first stable and clinically significant reduction in patient PTH for the vitamin D compound, and
- b. administering the initial dose of the vitamin D compound <u>determined in step a</u> to the patient.
- 8. (Original) The method of claim 7 wherein the vitamin D compound is paricalcitol.
- 9. (Original) The method of claim 8 wherein the initial dose is about <u>patient</u> <u>baseline parathyroid hormone/80 (bPTH/80)</u>.
- 10. (Amended) A method of treating a patient [undergoing vitamin D therapy] for end stage renal disease [ESRD] using a vitamin D therapy, [wherein the] comprising administering an initial dose of vitamin D [administered] to the patient wherein the initial dose of vitamin D is about patient baseline parathyroid hormone/80 (bPTH/80) and bPTH is the baseline PTH for the patient.
- 11. (Amended) A method of treating a patient [undergoing vitamin D therapy] for secondary hyperparathyroidism using a vitamin D therapy, [wherein the] comprising administering an initial dose of vitamin D [administered] to the patient wherein the initial dose of vitamin D is about patient baseline parathyroid hormone/80 (bPTH/80) and bPTH is the baseline PTH for the patient.
- 12. (Amended) A method of <u>determining the initial dose of a vitamin D</u> <u>compound</u> using a zero-intercept linear regression model [to determine the initial dose of a vitamin D compound].
- 13. (AMENDED) A method of treating a patient undergoing vitamin D therapy for [ESRD] end stage renal disease wherein a zero-intercept regression model is used to determine the initial dose of the vitamin D compound.
- 14. (Amended) The method of claim 13, wherein the vitamin D therapy [the vitamin D compound] results in the prevention or treatment of renal osteodystrophy or secondary hyperparathyroidism.
 - (Original) A method of claim 8 wherein the initial dose is at least 1 mcg.
 - 16. (New) The method of claim 13, wherein the vitamin D therapy does not increase the incidence of hypercalcemia.